## Copy for the Elected Office (EO/US'

# ATENT COOPERATION TREATY

	From the INTERNATIONAL BUREAU
PCT	То:
NOTIFICATION OF THE RECORDING OF A CHANGE  (PCT Rule 92bis.1 and Administrative Instructions, Section 422)  Date of mailing (day/month/year) 04 October 2000 (04.10.00)	KEITH W NASH & CO. 90-92 Regent Street Cambridge CB2 1DP ROYAUME-UNI
Applicant's or agent's file reference	INADODTANT MOTIFICATION
ML/C155.1/0	IMPORTANT NOTIFICATION
International application No. PCT/GB99/01650	International filing date (day/month/year) 26 May 1999 (26.05.99)
The following indications appeared on record concerning:      X the applicant      X the inventor	the agent the common representative    State of Nationality   State of Residence
Name and Address  LAMBERT, Peter, Anthony	GB GB
10 Dunton Close Sutton Coldfield Birmingham B75 5QD	Telephone No.
United Kingdom	Facsimile No.
	Teleprinter No.
2. The International Bureau hereby notifies the applicant that the	the following change has been recorded concerning:
the person the name X the add	
Name and Address	State of Nationality State of Residence  GB GB
LAMBERT, Peter, Anthony 125 Walsall Road Sutton Coldfield West Midlands B74 4NR	Telephone No.
United Kingdom	Facsimile No.
	Teleprinter No.
3. Further observations, if necessary:	
4. A copy of this notification has been sent to:	
X the receiving Office	the designated Offices concerned
the International Searching Authority	X the elected Offices concerned
the International Preliminary Examining Authority	other:
The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland	Authorized officer  Jocelyne Rey-Millet
Faccimile No. : (41, 22) 740, 14, 25	Telephone No : (41,-22) 338 83 38

# LATENT COOPERATION TREATY

To:

## **PCT**

### **NOTIFICATION OF ELECTION**

(PCT Rule 61.2)

LAMBERT, Peter, Anthony et al

Assistant Commissioner for Patents United States Patent and Trademark Office

Box PCT

Washington, D.C.20231 ÉTATS-UNIS D'AMÉRIQUE

Date of mailing (day/month/year) 15 December 1999 (15.12.99)	in its capacity as elected Office
International application No. PCT/GB99/01650	Applicant's or agent's file reference ML/C155.1/0
International filing date (day/month/year) 26 May 1999 (26.05.99)	Priority date (day/month/year) 28 May 1998 (28.05.98)
Applicant	

1.	The designated Office is hereby notified of its election made:
	X in the demand filed with the International Preliminary Examining Authority on:
	25 November 1999 (25.11.99)
	in a notice effecting later election filed with the International Bureau on:
	· · · · · · · · · · · · · · · · · · ·
2.	. The election X was
	was not
	made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).
	·

The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland **Authorized officer** 

Juan Cruz

Telephone No.: (41-22) 338.83.38

Facsimile No.: (41-22) 740.14.35



### INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference ML/C155.1/0	FOR FURTHER see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.  ACTION	
International application No.	International filing date (day/month/year)	(Earliest) Priority Date (day/month/year)
PCT/GB 99/01650	26/05/1999	28/05/1998
Applicant		
OXOID LIMITED et al.		
This International Search Report has bee according to Article 18. A copy is being tra	n prepared by this International Searching Aut ansmitted to the International Bureau.	thority and is transmitted to the applicant
This International Search Report consists  X  It is also accompanied by	of a total of sheets. a copy of each prior art document cited in this	s report.
Basis of the report		
a. With regard to the language, the language in which it was filed, un	international search was carried out on the balless otherwise indicated under this item.	asis of the international application in the
Authority (Rule 23.1(b)).	vas carried out on the basis of a translation of	
With regard to any nucleotide are     was carried out on the basis of th	nd/or amino acid sequence disclosed in the i	nternational application, the international search
	onal application in written form.	
filed together with the inte	ernational application in computer readable for	rm.
<u> </u>	this Authority in written form.	
1	this Authority in computer readble form.	
international application a	bsequently furnished written sequence listing or as filed has been furnished.	
the statement that the inf furnished	ormation recorded in computer readable form	is identical to the written sequence listing has been
2. Certain claims were fou	ind unsearchable (See Box I).	
3. Unity of invention is lac	king (see Box II).	
4. With regard to the title,		
1	ubmitted by the applicant.	
X the text has been establis ANTIBACTERIAL DIASCCH	shed by this Authority to read as follows:	
ANTIDACTENTAL DIAGOON	THE DENSTRICTS	
5. With regard to the abstract,	•	
1'	ubmitted by the applicant.	
the text has been established		rity as it appears in Box III. The applicant may, eport, submit comments to this Authority.
6. The figure of the drawings to be pub	lished with the abstract is Figure No.	2
X as suggested by the app	licant.	None of the figures.
because the applicant fai	iled to suggest a figure.	
because this figure bette	r characterizes the invention.	

Form PCT/ISA/210 (first sheet) (July 1998)

INTERNATIONAL SEARCH REPORT

PC1/G

Box III TEXT OF THE ABSTRACT (Continuation of item 5 of the first sheet)

Disclosed is an isolated compound having the structure shown in Figure 2, wherein n is an integer between 3 and 10 (inclusive) and X may be H, OH, alkyl, aryl, amyl, or an amino acid residue (optionally substituted) or a sugar residue (optionally substituted), and wherein K and K<sup>†</sup> are hydrophobic hydrocarbon or fatty acid chains (R may be the same as K<sup>†</sup> or different), a method of preparing compositions comprising the compound, and a diagnostic method comprising use of the composition.

Form PCT/ISA/210 (continuation of first sheet (2)) (July 1998)

# **PCT**

rec'd	1 8	AUG	2000	
WIPC	)		PCT	

# INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

MJL/C15	5.1/0	FOR FURTHER ACTION	Preliminary Examination Report (Form PCT/IPEA/416)
Internations	al application No.	International filing date (day/mont	h/year) Priority date (day/month/year)
PCT/GB9	99/01650	26/05/1999	28/05/1998
Applicant OXOID L  1. This is	IMITED et al.	nation report has been prepare	d by this International Preliminary Examining Authority
2. This F ⊠ TI be (s	REPORT consists of a total of his report is also accompanied and are the bas	5 sheets, including this cover so by ANNEXES, i.e. sheets of the standard sheets of the forthis report and/or sheets on the Administrative Instruction	ne description, claims and/or drawings which have
3. This re II III IV V VI VII VIII	<ul> <li>□ Lack of unity of invention</li> <li>☑ Reasoned statement uncitations and explanation</li> <li>□ Certain documents cited</li> <li>□ Certain defects in the interest</li> </ul>	pinion with regard to novelty, inv n der Article 35(2) with regard to ns suporting such statement d	rentive step and industrial applicability novelty, inventive step or industrial applicability;
Date of subn	nission of the demand	Date of d	completion of this report
25/11/199	9	16.08.20	00
preliminary e	ailing address of the international xamining authority:	Authorize	ed officer
<u></u>	European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 6 Fax: +49 89 2399 - 4465		r, S-E ne No. +49 89 2399 8554

Applicant's or agent's file reference

# INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB99/01650

#### I. Basis f the report

1. This report has been drawn on the basis of (substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments.): Description, pages: 1-34 as originally filed Claims, No.: 1-34 with telefax of 07/07/2000 Drawings, sheets: 1/8-8/8 as originally filed 2. The amendments have resulted in the cancellation of: ☐ the description, pages: ☐ the claims, Nos.: ☐ the drawings, sheets: 3. 

This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

4. Additional observations, if necessary:

# INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB99/01650

- V. Reasoned stat ment under Article 35(2) with r gard to novelty, inventiv step or industrial applicability; citations and explanations supporting such statement
- 1. Statement

Novelty (N)

Yes:

Claims 1-34

No:

Claims

Inventive step (IS)

Yes: No: Claims 1-34

140.

Claims

Industrial applicability (IA)

Yes:

Claims 1-34

No: Claims

2. Citations and explanations

see separate sheet

### VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether th claims are fully supported by the description, are made:

see separate sheet

## **EXAMINATION REPORT - SEPARATE SHEET**

#### V. Reasoned statement

The following documents will be referred to in this report:

D1 = Analytical Biochemistry; 1993, pages 49-56

D2 = Carbohydrate Research; 1984, pages 147-163

## Novelty (Article 33(2) PCT)

D1, which is referred to on page 3 (top), discloses i.a. peaks that may contain overlapping compounds.

Whether these - in such case - should be considered novelty-destroying will be a matter to settle under later national/regional regulations.

### Inventive step (Article 33(3) PCT)

No objection in view of the cited prior art.

### VIII. Certain observations

Claims:

Claim 2 has been added in view of D2, which discloses an accidentally overlapping compound 4 (n=3).

Since Claim 2 is dependent on Claim 1, the disclaimer should rather have been added to the independent claim.

[Whether the differing wordings of these claims, "isolated compound" and "compound", are of importance has to be settled in a later national/regional phase.]

2.

Figure 2 should preferably be included in Claim 1.

- 3. The definition "X may be" in Claim 1 is unclear.
- 4. Claims 5 and 8 should preferably refer to the compound of Claim 1 in order to avoid repeating the various definitions [subject to national requirements].
- 5. Claim 19 has been accepted because of the wording "specific binding".
- 6. The Applicant may later have to provide copies of the deposit receipts; see e.g. Rule 28 EPC.
- 7. Claims 30-31 and 33 include the step of administering a composition to a mammalian subject.

Such claims may not be acceptable under all national/regional regulations; in case of a later European phase the claims can be redrafted with a view to the Guidelines, C-IV, 4.2.

### **CLAIMS**

- 1. An isolated compound having the structure shown in Figure 2, wherein n is an integer between 3 and 10 (inclusive) and X may be H, OH, alkyl, aryl, amyl, or an amino acid residue (optionally substituted) or a sugar residue (optionally substituted), and wherein R and R<sup>1</sup> are hydrophobic hydrocarbon or fatty acid chains (R may be the same as R<sup>1</sup> or different).
- 2. A compound according to claim 1, where n is an integer other than 3.
- 3. A compound according to claim 1 or 2, wherein n = 6
- 4. A compound according to any one of claims 1, 2 or 3, wherein X = H, OH, D-alanyl or N-acetyl glucosamine.
- 5. A composition, comprising a compound in substantially pure form having the structure shown in Figure 2, wherein n is an integer between 3 and 10 (inclusive) and X is H, OH, alkyl, arryl, arryl, or an amino acid residue (optionally substituted) or a sugar residue (optionally substituted), and R and R<sup>1</sup> are hydrophobic hydrocarbon or fatty acid chains (R may be the same as R<sup>1</sup>, or different).
- 6. A composition according to claim 5, comprising an isolated compound in accordance with any one of claims 1 to 4.
- 7. A composition according to claim 5 or 6, in the form of a freeze-dried solid, an aqueous solution, or immobilised on a solid support.
- 8. A method of testing for a Gram +ve bacterial infection in a mammalian (typically, human) subject, the method comprising the steps of obtaining a sample of body fluid from the subject; contacting the sample with a composition comprising a compound having the structure shown in Figure 2, wherein n is an integer between 3 and 10 (inclusive) and X is H. OH, alkyl, aryl, aroyl, or an amino acid residue (optionally substituted) or a sugar

residue (optionally substituted), and R and R<sup>1</sup> are hydrophobic hydrocarbon or fatty acid chains (R may be the same as R<sup>1</sup>, or different); and detecting binding of antibodies (if any) in the sample to the composition.

- 9. A method according to claim 8, wherein the sample of body fluid obtained from the subject comprises whole blood, serum, urine or saliva.
- 10. A method according to claim 8 or 9, comprising the detection of binding to the composition of IgG antibodies in the sample.
- 11. A method according to any one of claims 8, 9 or 10, wherein the test method comprises the performance of an enzyme-linked immunosorbent assay (ELISA), radioimmunoassay (RIA), or a Western blot.
- 12. A method according to any one of claims 8 to 11, for testing for infection caused by Gram +ve cocci.
- 13. A method according to any one of claims 8 to 12, for testing for infection by a Streptococcus, a Staphylococcus or an Enterococcus.
- 14. A method according to any one of claims 8 to 13, for diagnosing the presence of a Gram +ve infection associated with a central venous catheter, a cerebrospinal fluid shunt or a prosthetic device.
- 15. A method according to any one of claims 8 to 14, wherein the composition is in accordance with any one of claims 5-7.
- 16. A diagnostic test kit for diagnosing the presence of a Gram +ve infection in a manimalian subject, the kit comprising: a solid support for performing a diagnostic test; and a composition in accordance with any one of claims 5-7.
- 17. A kit according to claim 16, further comprising one or more of the following: labelled

37

antibody; enzyme substrate; control sample; buffer, and instructions for use.

- 18. A sterile vaccine composition for use against a Gram +ve infection in a mammalian subject, the vaccine comprising an isolated compound in accordance with any one of claims 1 to 4, or a composition in accordance with any one of claims 5 to 7.
- 19. An isolated immunoglobulin molecule or variant thereof having specific binding for a compound in accordance with any one of claims 1 to 4.
- 20. An isolated eukaryotic cell producing an immunoglobulin molecule or variant thereof in accordance with claim 19.
- 21. A method of making a composition in accordance with any one of claims 5 to 7, the method comprising the steps of: culturing a Gram +ve bacterium in a growth medium so as to cause the bacterium to secrete into the growth medium the compound having the structure shown in Figure 2; separating the growth medium from the bacterial cells; fractionating the growth medium; and isolating that fraction which comprises, in substantially pure form, the compound having the structure shown in Figure 2.
- 22. Staphylococcus epidermidis strain CAN 6KIII, deposited under accession number NCIMB 40896.
- 23. Staphylococcus epidermidis strain HAR 6KIV, deposited under accession number NCIMB 40945.
- 24. Staphylococcus epidermidis strain COS 6KV, deposited under accession number NCIMB 40946.
- 25. Staphylococcus epidermidis strain MIL 6LI, deposited under accession number NCIMB 40947.
- 26. Staphylococcus epidermidis strain HED 6LL, deposited under accession number

38

NCIMB 40948.

- 27. Staphylococcus haemolyticus strain ONE 6KVI, deposited under accession number NCIMB 40949.
- 28. Micrococcus kristinae strain MAT 6LII, deposited under accession number NCIMB 40950.
- 29. A method according to claim 21, comprising the step of culturing one or more organisms selected from the group consisting of: Staphylococcus epidermidis strain CAN 6KIII; Staphylococcus epidermidis strain HAR 6KIV; Staphylococcus epidermidis strain COS 6KV; Staphylococcus epidermidis strain MIL 6LI; Staphylococcus epidermidis strain HED 6LI; Staphylococcus haemolyticus strain ONE 6KVI; Micrococcus kristinae strain MAT 6LII.
- 30. A method of making an immunoglobulin having specific binding for a melecule in accordance with claim 1, the method comprising the steps of: preparing a composition comprising a compound in accordance with any one of claims 1-4; administering the composition to a mammalian subject; and obtaining from the subject a sample comprising antibodies or antibody-producing cells.
- 31. A method according to claim 30, wherein antibody-producing cells are isolated from the subject and used to prepare a hybridoma.
- 32. A method of obtaining an immunoglobulin or antigen-binding variant thereof having specific binding for a compound in accordance with any one of claims 1-4, the method comprising the steps of: screening a library of viruses or other particles displacing an immunoglobulin or antigen-binding variant thereof on their surface; and selecting those members of the library which display an immunoglobulin or antigen-binding variant thereof which bind to the compound.
- 33. A method of inducing antibodies in a human subject, the method comprising the steps

39

of preparing a physiologically acceptable composition in accordance with claim 5; and administering the composition to the subject.

34. A vaccine for inducing antibodies in a mammalian subject the vaccine comprising a composition in accordance with claim 5 and a physiologically acceptable excipient, carrier or diluent.

# (19) World Intellectual Prop rty Organization International Bureau



# 

# (43) International Publication Date 2. December 1999 (02.12.1999)

## **PCT**

# (10) International Publication Number WO 99/61913 A3

(51) International Patent Classification<sup>6</sup>: C07H 3/04, C07K 1/00, A61K 31/70, C12N 15/06, 1/20, A61K 39/085, C07K 16/12 // (C12N 1/20, C12R 1:45)

West Midlands B74 4NR (GB). ELLIOTT, Thomas, Stuart, Jackson [GB/GB]; 57 Roman Lane, Little Aston Park, Sutton Coldfield, Birmingham B74 3AE (GB).

- (21) International Application Number: PCT/GB99/01650
- (74) Agent: KEITH W NASH & CO.; 90-92 Regent Street, Cambridge CB2 1DP (GB).

(84) Designated States (regional): European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC,

- (22) International Filing Date: 26 May 1999 (26.05.1999)
- (81) Designated State (national): US.

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

9811347.5

28 May 1998 (28.05.1998) G

Published:

NL, PT, SE).

- With international search report.
- (71) Applicant (for all designated States except US): OXOID LIMITED [GB/GB]; Kingsland Business Park, Wade Road, Basingstoke, Hampshire RG24 8PW (GB).
- (88) Date of publication of the international search report: 19 April 2001

- (72) Inventors; and
- (75) Inventors/Applicants (for US only): LAMBERT, Peter, Anthony [GB/GB]; 125 Walsall Road, Sutton Coldfield,

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

### (54) Title: ANTIBACTERIALS DIASCCHARIDE DERIVATIVES

$$H = \begin{bmatrix} -0 - CH_2 \\ X + \\ CH_2 - 0 - P - \\ 0 \end{bmatrix}_{n} = \begin{bmatrix} -0 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH \\ OH \\ OH \end{bmatrix}_{OH} = \begin{bmatrix} -CH_2 - CH_2 \\ OH$$

(57) Abstract: Disclosed is an isolated compound having the structure shown in Figure 2, wherein n is an integer between 3 and 10 (inclusive) and X may be H, OH, alkyl, aryl, amyl, or an amino acid residue (optionally substituted) or a sugar residue (optionally substituted), and wherein R and R<sup>1</sup> are hydrophobic hydrocarbon or fatty acid chains (R may be the same as R<sup>1</sup> or different), a method of preparing compositions comprising the compound, and a diagnostic method comprising use of the composition.

A. CLASSIFICATION OF SUBJECT MATTER IPC 6 C07H3/04 C07K1/00 A61K39/085

C07K16/12

A61K31/70 C12N15/06 //(C12N1/20,C12R1:45)

C12N1/20

According to International Patent Classification (IPC) or to both national classification and IPC

### B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols) IPC 6 CO7H CO7K A61K C12N

Documentation searched other than minimum documentation to the extent that such documents are included. In the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS	CONSIDERED TO BE	E RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	J.J.OLTVOORT ET AL.: "Synthesis of a Lipoteichoic Acid-Carrier Fragment of Staphylococcus aureus." CARBOHYDRATE RESEARCH., vol. 130, 1984, pages 147-163, XP002120748 ELSEVIER SCIENTIFIC PUBLISHING COMPANY. AMSTERDAM., NL ISSN: 0008-6215	1,4,7
A	page 148, compound 4 WO 97 42343 A (PHARMACIA AND UPJOHN COMPANY) 13 November 1997 (1997-11-13) the whole document/	1,4,7, 15,18-20

Į	Special categories of cited documents :	
	"A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international	T later document published after the international fiting date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
I	ming date	"X" document of particular relevance; the claimed invention
ı	"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	Involve an inventive step when the document is taken alone document of partial ar releasement the stellar of th
ł	of document referring to an oral disclosure, use, exhibition or other means	document is combined with one or man other the
l	*P* document published prior to the international filing date but later than the priority date claimed	In the art.
Г	Data of the	"&" document member of the same patent family

Date of the actual completion of the international search

Date of mailing of the international search report

# 28 October 1999

Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016

Further documents are listed in the continuation of box C.

11/11/1999 Authorized officer

Scott, J

Patent family members are listed in annex.

C.(Continu	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	PC1/GB 99/01650
Category *	Citation of document, with Indication, where appropriate, of the relevant passages	Relevant to claim No.
A	W.FISCHER: "Molecular Analysis of Lipid Macroamphiphiles by Hydrophobic Interaction Chromatography Exemplified with Lipotechoic Acids." ANALYTICAL CHEMISTRY, vol. 208, 1993, pages 49-56, XP002120749 cited in the application abstract	1,4
A	H.I.WERGELAND ET AL: "Antibodies to Staphylococcal Peptidoglycan and its Peptide Epitopes, Techoic Acid, and Lipotechoic Acid in Sera from Blood Donors and Patients with Staphylococcal Infections."  JOURNAL OF CLINICAL MICROBIOLOGY, vol. 27, no. 6, June 1989 (1989-06), pages 1286-1291, XP002120750 cited in the application abstract	1,4
A	J.J.OLTVOORT ET AL.: "A Simple Approach to the Synthesis of a Membrane Techoic Acid Fragment of Staphylococcus aureus." RECUEIL, JOURNAL OF THE ROYAL NETHERLANDS CHEMICAL SOCIETY, vol. 101, no. 3, March 1982 (1982-03), pages 87-91, XP002120751 the whole document	1,4
	DATABASE WPI Section Ch, Week 198703 Derwent Publications Ltd., London, GB; Class B04, AN 1987-017786 XP002120752 & JP 61 275217 A (YAKULT HONSHA KK), 5 December 1986 (1986-12-05) abstract	1,4



Int tal Application No
PCT/GB 99/01650

Patent document cited in search report	ł	Publication date	Patent family member(s)	Publication date
WO 9742343	A	13-11-1997	AU 2746997 A EP 0901529 A WO 9742311 A	26-11-1997 17-03-1999 13-11-1997
JP 61275217	A	05-12-1986	NONE	